UNITED STATES DISTRICT COURT NORTHERN DISTRICT OF CALIFORNIA

ORACLE CORPORATION, ET AL.,

No. C 11-00910 JCS

Plaintiffs,

v.

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DRUGLOGIC, INC.,

Defendant.

ORDER GRANTING IN PART AND DENYING IN PART PLAINTIFFS⁷ MOTION TO DISMISS AND STRIKE DRUGLOGIC'S SECOND AMENDED DEFENSES AND COUNTERCLAIMS [Docket No. 79]

INTRODUCTION I.

Plaintiffs Oracle Corporation and Oracle International Corporation ("Oracle") filed this action against Defendant DrugLogic, Inc. ("DrugLogic") alleging infringement by DrugLogic of Oracle's U.S. Patent No. 6,684,221 ("the '221 patent") and seeking a declaratory judgment of noninfringement and invalidity as to DrugLogic's U.S. Patent No. 6,789,091 ("the '091 patent"). In response, DrugLogic asserted various affirmative defenses and counterclaims, the sufficiency of which Oracle challenged in a motion to dismiss and strike filed in May 2011. The Court granted in part and denied in part Oracle's motion, dismissing certain counterclaims and affirmative defenses with leave to amend. See Docket No. 54 ("the August 8 Order"). Presently before the Court is Oracle's Motion to Dismiss and Strike Druglogic's Second Amended Defenses and Counterclaims ("the Motion"). The Court finds the Motion appropriate for disposition without oral argument pursuant to Civil Local Rule 7-1(b). For the reasons stated below, the Motion is GRANTED in part and DENIED in part.

II. BACKGROUND¹

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A. The Court's August 8, 2011 Order

In its August 8 Order, the Court addressed the adequacy of various affirmative defenses and counterclaims asserted by DrugLogic in its original answer. With respect to DrugLogic's inequitable conduct affirmative defense and counterclaim, which was based on the theory that Oracle had withheld material information from the patent examiner during the application process, the Court found DrugLogic's allegations were insufficient under Exergen Corp. v. Wal-Mart Stores, Inc., 575 F.3d 1312 (Fed. Cir. 2009) and Federal Rule of Civil Procedure 9(b). August 8 Order at 17. In particular, the Court held that DrugLogic had not sufficiently alleged the "what" and "where" of the alleged inequitable conduct, stating that "[allthough DrugLogic identifies potentially material information contained in the allegedly withheld references by noting that WHO-Drug, COSTART, Read Codes, CPT, Unified Medical Language System, Metathesaurus, MeSH, and PubMed are all 'hierarchical relational medical thesauruses,' some of which contain 'clinical terms used in conjunction with clinical studies,' DrugLogic fails to allege where specifically in those references that material information could be found." *Id.* The Court went on to hold that DrugLogic had not sufficiently alleged the "why" and "how" of its inequitable conduct defense. *Id.* at 18. In particular, the Court found that DrugLogic had not alleged "any facts to support an inference that the information allegedly withheld from the PTO is not cumulative of other information previously disclosed to the examiner, particularly given that Oracle referenced WHO-Drug, COSTART, and CPT in the '221 patent specification." *Id.* As to the question of whether DrugLogic's allegations were sufficient to give rise to an inference of deceptive intent, the Court held: "[A]lthough this is a close question, the Court finds that if DrugLogic has adequately pled knowledge as described above, no additional pleading will be necessary with respect to deceptive intent." *Id.* at 19. The Court dismissed the counterclaim and struck the affirmative defense based on inequitable conduct and granted DrugLogic leave to amend its inequitable conduct allegations. *Id.* at 25.

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¹A more detailed overview of the procedural background of the case is included in the Court's August 8, 2011 Order.

B. DrugLogic's Second Amended Answer²

In its Second Amended Answer, Defenses, Counterclaims and Demand for Jury Trial of Defendant and Counterclaimant DrugLogic, Inc. ("SAA"), Druglogic offers more detailed allegations regarding the patentee's alleged failure to disclose material information to the patent examiner and asserts a number of new state law counterclaims, including a counterclaim for unfair competition.

DrugLogic's amended inequitable conduct allegations including the following:

- 34. The provisional application from which the '221 Patent claims priority, U.S. Provisional Application No. 60/132,926, filed by inventor Kim Rejndrup and his patent counsel Rodney Johnson, identifies MedDRA, WHO-Drug, WHO-Art, ICD9, ICD10 and COSTART as "hierarchical structured dictionaries."
- 35. MedDRA, WHO-Art and COSTART were and are all hierarchical relational medical thesauruses, and all of them contain clinical terms that are or have been used in or derived from clinical studies. Inventor Kim Rejndrup and his patent attorneys Rodney Johnson and Christopher Lutz knew that MedDRA, WHO-Art and COSTART were and are hierarchical relational medical thesauruses, and that they contain clinical terms that are or have been used in or derived from clinical studies.

. . .

- 44. The information withheld -- that WHO-Art, COSTART, and MedDRA were and are hierarchical relational medical thesauruses that contain clinical terms used in conjunction with or derived from clinical studies -- is relevant to claims 1, 21, 51 and 53, and in particular to the limitations of those claims that require (i) defining (or identifying) a plurality of clinical terms for a clinical study and (ii) storing the plurality of terms in a memory according to a hierarchy of relations. WHO-Art, COSTART, and MedDRA are medical thesauruses, and each of them contain clinical terms used in conjunction with or derived from clinical studies. In order for these clinical terms used in conjunction with or derived from clinical studies to have been entered into WHO-Art, COSTART, and MedDRA, the clinical terms for clinical studies were first identified and then stored in a memory according to a hierarchy of relations.
- 45. The withheld information -- that WHO-Art, COSTART, and MedDRA are hierarchical relational medical thesauruses that contain clinical terms used in conjunction with or derived from clinical studies is present at, and can be found in, any portion of WHO-Art, COSTART, and MedDRA. That is, just as the fact that Roget's Thesaurus is a thesaurus containing English-language words can be ascertained from virtually any page in Roget's Thesaurus, the fact that WHO-Art, COSTART and MedDRA are hierarchical relational medical thesauruses containing clinical terms used in or derived from clinical studies would

²On September 2, 2011, DrugLogic filed a First Amended Answer. *See* Docket No. 71. Following a meet and confer, the parties stipulated to the filing of a Second Amended Answer to address certain concerns expressed by Oracle relating to the sufficiency of the amended counterclaims and defenses. *See* Docket No. 75. The Second Amended Answer was filed on September 23, 2011.

be immediately perceived by someone reviewing any portion of the contents of WHO-Art, COSTART or MedDRA. For example, if one were to look up in MedDRA the clinical term "cardiac flutter," a term associated with certain clinical studies, one would find that the term is categorized in a hierarchical structure, with synonyms (reflecting that MedDRA is a thesaurus) and related terms having the same level of detail or specificity categorized at the same level, and with related terms having broader meanings (reflecting that MedDRA is relational) categorized at correspondingly "higher" levels (reflecting that MedDRA is hierarchical). One would find similar clinical terms and organization with COSTART and WHO-ART. Inventor Kim Rejndrup and his patent lawyers were aware of the content and organization of WHO-Art, COSTART and MedDRA, specifically that they were relational medical thesauruses containing clinical terms, and they knew that this information was material to the claims of the '221 Patent.

- 46. The information withheld is material because if the patent examiner had been aware of the actual nature of the withheld information, that is, that WHO-Art, COSTART, and MedDRA are each a hierarchical relational medical thesaurus that contains clinical terms used in conjunction with or derived from clinical studies, the examiner would not have allowed the claims to issue in their present form. Instead, the examiner would have found that the withheld information provides a teaching that was absent from the prior art relied upon by the examiner, and he would have rejected the claims instead of allowing them.
- 47. Specifically, the examiner allowed the claims only after the words "clinical terms" and "clinical studies" were added to the claims. If the examiner had been told that WHO-Art, COSTART, and MedDRA were medical thesauruses that contain clinical terms from clinical studies, the examiner would have maintained the rejections and not allowed the claims to issue.
- 48. The information withheld is not cumulative to other information provided to the examiner because neither inventor Kim Rejndrup nor his patent attorneys Rodney Johnson and Christopher Lutz provided any prior art to the USPTO, and because the incomplete disclosure of WHO-Drug, COSTART and CPT in the specification of the '221 Patent fails to make any reference to WHO-Art or to MedDRA, and it fails to state that any of them are thesauruses of medical or clinical terms. The incomplete description and disclosure of the prior art materially misrepresents its nature and scope and fails to put the examiner on notice of its relevance. Indeed, it undoubtedly led the examiner to the conclusion that this prior art was not relevant.
- 49. The information withheld is not cumulative to other information cited by the examiner because none of the prior art cited by the examiner was or disclosed a medical thesaurus containing clinical terms, in which medical (including clinical) terms are stored according to a hierarchy of relations, including relations indicative of associations between medical (including clinical) terms.
- 50. On information and belief, the failure of inventor Kim Rejndrup and his patent attorneys Rodney Johnson and Christopher Lutz, to provide copies of or excerpts from one or more of the WHO-Art, COSTART, and MedDRA hierarchical relational medical thesauruses to the U.S. Patent and Trademark Office, and their failure to advise the U.S. Patent and Trademark Office that WHO-Art, COSTART, and MedDRA contain clinical terms used in conjunction with or derived from clinical studies, were intentional acts or omissions, done with deceptive intent.

SAA, ¶¶ 34-35, 44-50.

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C. The Motion

In the Motion, Oracle argues that: 1) Druglogic's counterclaim and affirmative defense of inequitable conduct should be dismissed (as to the counterclaim) and stricken (as to the affirmative defense) on the basis that the specific allegations addressing what was not disclosed to the Patent and Trade Office ("PTO") show that Oracle did not, in fact, withhold any material information from the PTO and therefore, the allegations do not give rise to an inference of deceptive intent; 2) DrugLogic's allegation of common law unfair competition in its Sixth Claim for Relief should be dismissed because under Bank of the West v. Superior Court, 2 Cal. 4th 1254 (1992), unfair competition claims asserted under common law are limited to the act of "passing off," and no passing off has been alleged here; and 3) DrugLogic's requests for damages on its unfair competition counterclaim and for injunctive relief on its breach of contract counterclaim should be stricken because these remedies are not available on DrugLogic's claims. Motion at 2-3.

1. **Inequitable Conduct Allegations**

Oracle argues that the new details added to DrugLogic's allegations do not save the defense and counterclaim but instead reveal that they are factually implausible under the standard set forth in Bell Atlantic Corp. v. Twombly, 550 U.S. 544, 555 (2007) and Ashcroft v. Iqbal, 129 S. Ct. 1937, 1950 (2009). *Id.* at 5. According to Oracle, DrugLogic's theory – that the inventor and others who prosecuted the patent hid from the Patent Office the fact that prior art systems described in the '221 specification "were and are all hierarchical relational medical thesauruses ... [that] contain clinical terms that are or have been used in or derived from clinical studies" – is demonstrably false on the face of the '221 patent because it is clear from the specification that all of the prior art references are: 1) medical thesauruses; 2) "hierarchical"; and 3) relational. Id. It is also clear from the specification, Oracle argues, that they contain clinical terms derived from clinical studies. *Id.*

To show that it would have been apparent to the patent examiner that the systems described in the prior art referenced in the '221 patent were medical thesauruses, Oracle points to the titles of the systems used in the specification. Id. (citing '221 Pat., col. 4, lines 4-10 ("Common vendorsupplied dictionaries include WHO-Drug (World Health Oragnization Drug Dictionary) by the

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World Health Organization, COSTART (Coding Symbols for a Thesaurus of Adverse Reaction
Terms) by the Drug Information Association, and CPT (Current Procedural Terminology) by the
American Medical Association")). Id. Oracle rejects DrugLogic's suggestion that the nature of the
systems referenced in the specification would not have been apparent to the patent examiner because
they were referred to as dictionaries rather than thesauruses, id . (citing SAA ¶¶ 34-35), pointing out
that one of the systems contains the word "thesaurus" in its title; to the extent that the other systems
are referred to as dictionaries, Oracle asserts, this is an accurate description of these systems as they
contain the word "dictionary" in their titles, and therefore, no deceptive intent can be inferred from
referring to this prior art as such. <i>Id</i> .

It is also clear from the face of the specification that the prior art systems were hierarchical, Oracle argues. *Id.* at 7. Oracle points to Figures 1 and 2 and the accompanying text in the specification that describes these figures in support of its position. *Id.* (citing '221 Pat., col. 4, lines 12-16 ("Fig. 1 shows the various entities that interact with the classification and mapping system, and Fig. 2 shows the structure of a vendor-supplied dictionary used with such a system. Referring to Figs. 1 and 2, the hierarchical structure of the WHO-Drug dictionary 44 is shown").

Oracle further contends that it is was apparent from the specification that the prior art systems contained clinical terms that are or have been used in or derived from clinical studies. *Id.* In particular, Oracle argues that the "patent explicitly states that the purpose of the prior art systems in the claimed invention is to load clinical terms into that invention [and therefore] those systems must contain clinical terms." *Id.* (citing '221 Pat., col 3, lines 40-43 ("The *clinical terms* are initially transmitted to the thesaurus database 18 from an external media source, such as a CD-ROM 40, from a loader 42""); col. 4, lines 16-18 (*"Clinical terms* are read from the CD-ROM 40 and stored in the content table 20"); col. 5, lines 10-12 ("Referring to Figs. 1 and 5, a dictionary of *clinical terms* is loaded into the thesaurus database 18") (emphasis added in Motion)).

Finally, Oracle argues that the '221 specification disclosed that the prior art systems were relational, quoting the following language in the specification:

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FIG. 1 shows the various entities that interact with the classification and mapping system, and FIG. 2 shows the structure of a vendor supplied dictionary used with such a 15 system. Referring to FIGS. 1 and 2, the hierarchical structure of the WHO-Drug dictionary 44 is shown. Clinical terms are read from the CD-ROM 40 and stored in the content table 20. Relations between the clinical terms are also read and stored in the relation table 22 to define relations between terms on different levels 46, as defined by the hierarchy. In this manner, both the content and the hierarchical structure of the dictionary 44 are transmitted to the thesaurus database 18 via the loader 42 and stored in the content table 20 and the relation table 22.

Id. (quoting '221 Pat., col. 4, lines 12-24)(emphasis added in Motion).

Because all of the allegedly withheld information was in fact disclosed to the patent examiner in the specification, Oracle argues, DrugLogic has failed to allege a material omission or misrepresentation; therefore, its inequitable conduct claim should be dismissed. *Id.* at 9 (citing Funai Elec. Co. v. Daewoo Elecs. Corp., 2006 WL 3780715, at * 3-4 (N.D. Cal. Dec. 20, 2006), Chiron Corp. v. Genentech, Inc., 286 F. Supp. 2d 1126, 1137 (E.D. Cal. 2002)). Oracle further asserts that in the absence of any factual allegations showing a material misrepresentation or omission, DrugLogic's allegations also do not support a plausible inference of intent to deceive. *Id.* Oracle points out that under Exergen, to state a claim for inequitable conduct, DrugLogic's factual allegations must "plausibly suggest '[a] deliberate decision to withhold a known material reference or to make a knowingly false misrepresentation." *Id* (quoting *Exergen*, 575 F.3d at 1331 (internal quotations omitted)).

2. **Unfair Competition Counterclaim**

Oracle argues that DrugLogic has improperly combined in Counterclaim Six a statutory unfair competition claim under Cal. Bus. & Prof. Code §§ 17200 et seq. with a common law unfair competition claim. Id. at 10. The latter claim fails, Oracle asserts, because the California Supreme Court has limited unfair competition claims under common law to the act of "passing off" one's goods as those of another. Id. (citing Bank of the West v. Superior Court, 2 Cal. 4th 1254, 1263 (1992)). Oracle contends that because there is no allegation that Relsys or Oracle engaged in "passing off" their goods as those of DrugLogic, DrugLogic's claim for common law unfair competition should be dismissed.

3. Improper Remedies

a. Injunctive Relief for Breach of Contract

Oracle argues that DrugLogic's requested relief on Counterclaim II, for breach of contract, improperly includes a request for injunctive relief. *Id.* at 11 (citing SAA Counterclaim II Requested Relief, ¶ B ("seeking "[a]n injunction prohibiting Oracle...from further use of DrugLogic's confidential information and the software created through Relsys's impropert reverse engineering and decompiling")). According to Oracle, injunctive relief is available on a breach of contract claim only where a plaintiff adequately alleges entitlement to specific performance under the contract. *Id.* (citing Cal Civ. Code Section 3423(e) ("[a]n injunction may not be granted . . . [t]o prevent the breach of a contract the performance of which would not be specifically enforced"), *Golden West Baseball Co. v. City of Annaheim*, 25 Cal. App. 4th (1994) ("[a]n injunction to enforce the terms of a contract may only be issued if the contract is specifically enforceable")). Because DrugLogic has not alleged specific facts showing that it is entitled to specific enforcement and in particular, has not alleged any ongoing conduct that would be subject to specific performance, Oracle contends that this relief should be stricken from DrugLogic's breach of contract counterclaim. *Id.* (citing *Tamarind Lithography Workshop, Inc. v. Sanders*, 143 Cal. App. 3d 571, 575 (1983) for elements required for specific performance).

b. Compensatory Damages for Unfair Competition

Oracle argues that the Court should strike DrugLogic's request for "an award disgorging any benefit received by Oracle as a result of its unfair competition" as a remedy on Counterclaim VI, for unfair competition. *Id.* at 12. Oracle argues that California's statutory unfair competition law ("UCL"), only authorizes "two forms of equitable relief: *preventive.*, i.e., an injunction, and *restorative*, ie., an order for restitution." *Id.* (citing *Silvaco Data Sys. v. Intel Corp.*, 184 Cal. App. 4th 210, 244 (2010) and Cal. Bus. & Prof. Code Section 17203). The UCL does not authorize compensatory relief, Oracle contends. *Id.* Moreover, Oracle argues, California courts have expressly limited "restitution" to "orders compelling a [Section 17200] defendant to return money obtained through an unfair business practice to those persons in interest from whom the property

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was taken, that is, to persons who had an ownership interest in the property or those claiming through that person." Id. (quoting Krause v. Trinity Mgmt. Servs., Inc., 23 Cal. 4th 116, 126-27 (2000)). Because the monetary relief requested by DrugLogic falls outside the scope of the monetary relief available under the UCL, Oracle contends, the request for disgorgement should be stricken.

C. **Opposition**

In its Opposition, DrugLogic contends that: 1) its amended inequitable conduct allegations are sufficient to remedy the deficiencies identified by the Court in its August 8 Order; 2) its unfair competition claim does not fail to the extent it is based on common law because the Bank of the West case does not apply; 3) DrugLogic's request for an order disgorging benefits under its unfair competition claim does not fail because it is based on its common law unfair competition claim and not its statutory claim under Cal. Bus. & Profs. Code §§ 7200 et seq.; and 4) DrugLogic's request for an injunction under its claim for breach of contract is proper.

1. **Inequitable Conduct Allegations**

DrugLogic asserts that inequitable conduct is adequately alleged because DrugLogic has included in the SAA detailed allegations showing that the patentees misled the examiner by mischaracterizing the prior art referenced in the specification and failing to provide copies of the prior art to the examiner, which would have revealed to the examiner that any one of the prior art references disclosed the full scope of the invention claimed in the '221 patent. Opposition at 2, 5 (citing Advanced Ion Beam Tech., Inc. v. Varian Semiconductor Equip. Assoc., 721 F. Supp. 2d 62, 79 (D. Mass. 2010); Golden Valley Microwave Foods, Inc. v. Weaver Popcorn Co., Inc., 837 F. Supp. 1444 (N.D. In. 1992)). According to DrugLogic, Oracle mischaracterized the prior art at issue by failing to disclose to the examiner that "each of [th]e prior art references contain[ed] each of the four elements of the invention," namely 1) a thesaurus, 2) a hierarchy, 3) relations, and 4)clinical terms. *Id.* at 7 (emphasis in original). DrugLogic further points to the Manual of Patent Examining Procedures ("MPEP"), which requires that inventors and their attorneys disclose information material to patentability, include such information in their information disclosure statement and

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provide a "legible copy" of each publication listed in the information disclosure statement. *Id.* at 8 (citing MPEP § 1.56 and 37 C.F.R. 1.98).

DrugLogic rejects Oracle's assertion that it fully disclosed in the specification that the prior art at issue contained the elements of the claimed inventions, arguing that "these references to the prior art were scattered throughout the specification in an attempt to hide the fact that the invention claimed in the '221 Patent was unpatentable." *Id.* at 9. "More importantly," DrugLogic asserts, "even when viewed together, the information contained in the specification does not disclose the true nature and scope of each piece of prior art." *Id.* This omission is particularly significant, DrugLogic contends, because the examiner initially rejected each of the claims and only allowed the '221 Patent to issue when the limitation of clinical terms was added. *Id.* at 9-10 & Ex. A (Notice of Allowability).

With respect to Oracle's argument that DrugLogic has not alleged facts sufficient to show deceptive intent, DrugLogic contends that the Court already ruled on this question when it found that so long as DrugLogic adequately alleged knowledge of the prior art at issue by those who had a duty of disclosure, deceptive intent would also be adequately alleged. *Id.* at 5-6 (quoting August 8 Order at 18-19 ("the Court finds that if DrugLogic adequately pleads knowledge as described above, no additional pleading will be necessary with respect to deceptive intent under Exergen")).

DrugLogic rejects Oracle's reliance on Funai Elec. Co. v. Daewoo Elecs. Corp., 2006 WL 3780715 at * 3-4 (N.D. Cal. Dec. 20, 2006), arguing that that case is factually distinguishable because the material aspects of the prior art that the applicant allegedly failed to disclose to the examiner – a Japanese patent application – were, in fact, disclosed in an English translation of the Japanese Patent Office's evaluation of the application, which was provided to the Examiner. *Id.* at 10. DrugLogic also contends that Oracle's reliance on Chiron Corp. v. Genentech, Inc., 268 F. Supp. 2d 1126, 1137 (E.D. Cal. 2002) is misplaced. *Id.* at 11. According to DrugLogic, that case is not on point because it was decided after discovery and full development of the record, at which point the court concluded that there was no evidence that the information at issue was "intentionally obscured." Id. at 11.

2. Unfair Competition Counterclaim

DrugLogic rejets Oracle's assertion that under *Bank of the West*, common law unfair competition claims are limited to claims for "passing off." *Id.* at 12. According to DrugLogic, the central holding of the *Bank of the West* decision was that an insurance policy did not cover claims for advertising injury under California's Unfair Business Practices Act, Cal. Bus. & Prof. Code §§ 17200 *et seq.*, and therefore, the statements in that case about common law unfair competition claims were dicta. *Id.* DrugLogic cites to a decision by this Court, *Hewlett-Packard Co. v. Cigna Prop. & Cas. Ins. Co.*, 1999 U.S. Dist. LEXIS 20655 (N.D. Cal. Aug. 24, 1999), in which the court held that in the context of an insurance coverage dispute, claims for "unfair competition" would be understood by a lay person to include claims not only for "passing off" but also for false advertising and other tortious conduct. DrugLogic also points to a decision by the Ninth Circuit issued after the *Bank of the West, Duncan v. Stuezle*, 76 F.3d 1480 (9th Cir. 1996). According to DrugLogic, in this case the Ninth Circuit held "that a plaintiff properly asserted a claim for common law unfair competition where he claimed that the defendant misappropriated certain proprietary information regarding, among other things, marketing strategy, revenue and methods by which a particular product was produced." *Id.* at 13.

3. Improper Remedies

a. Compensatory Damages for Unfair Competition

DrugLogic does not dispute that compensatory damages are unavailable under the UCL and therefore, that disgorgement of profits is not an available remedy under the UCL. *Id.* It argues, however, that this remedy is available under its California's common law unfair competition claim, which survives for the reasons stated above. *Id* (citing *Duncan*, 76 F.3d at 1489-1490).

b. Injunctive Relief for Breach of Contract

DrugLogic argues that it has adequately alleged that pecuniary compensation is not adequate to remedy Oracle's alleged breach of contract and therefore, its request for an injunction under this claim is proper. *Id.* at 14. In particular, DrugLogic points to a number of allegations in its SAA,

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including its allegations that Oracle "continues to make, import, use, sell, and offer to sell Argus Perceptive (the improperly reverse engineered product) to customers . . ., which is a violation of the prohibition in the Co-Marketing Agreement from selling, transferring, publishing, disclosing or otherwise making available any portion of DrugLogic's confidential information to a third party." *Id.* at 14-15 (citing SAA, ¶¶103, 119, 121, 125 and p. 29 ¶ B). Based on these allegations, DrugLogics argues that it is merely requesting that "Oracle be compelled to specifically perform its continuing obligations under the Co-Marketing Agreement to refrain from certain activities." *Id.* at 15.

D. Reply

In its Reply brief, Oracle reiterates its argument that DrugLogic's allegations of inequitable conduct fail because they are implausible under Twombley and Iqbal in light of the disclosures in the specification. Reply at 2-3. Oracle rejects as "demonstrably wrong" DrugLogic's assertion that the disclosures in the specification are "scattered" throughout the specification and further asserts that the specification "makes clear that its description of each alleged prior art system applies to all of the alleged prior art systems." *Id.* at 4-6. Oracle also argues that the cases cited by DrugLogic – Advanced Ion Beam and Golden Valley – do not support DrugLogic's position. Id. at 7-8. Oracle argues that reliance on the patent prosecution guidelines in the MPEP misses the point because while those guidelines set forth the types of information an applicant is required to disclose, they do not address the types of facts that are sufficient to give rise to a reasonable inference that the applicant intended to deceive the patent examiner. Id. at 8. Finally, as to the intent to deceive, Oracle rejects DrugLogic's contention that this question has already been decided by the Court. Id. at 2. Rather, Oracle argues that the Court's holding regarding deceptive intent simply meant that "if DrugLogic adequately pled that the patent applicants knew about and withheld certain specific pieces of information about the allegedly withheld prior art, then they would also have adequately pled deceptive intent." Id. at 2.

As to the common law claim for unfair competition, Oracle argues neither Hewlett-Packard Co. v. Cigna Prop. Cas. Ins. Co. nor Duncan v. Stuetzle – the two cases cited by DrugLogic in

support of its contention that it has stated a claim for common law unfair competition – redefined common law unfair competition as set forth in Bank of the West. Id. at 9. Oracle also points out that DrugLogic does not dispute that compensatory damages are not available for statutory unfair competition under Cal. Bus. & Prof. Code §§ 17200 and therefore, to the extent the common law unfair competition counterclaim fails, so too does DrugLogic's request for disgorgement of profits on its unfair competition counterclaim.

Finally, as to the request for injunctive relief on DrugLogic's breach of contract claim, Oracle asserts that DrugLogic has not made any attempt to allege facts showing that the requirements for specific performance have been alleged. *Id.* at 10.

III. **ANALYSIS**

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Legal Standard³ A.

1. **Motion to Dismiss**

Federal Rule of Civil Procedure 8(a)(2) provides that a pleading must contain a "short and plain statement of the claim showing that the pleader is entitled to relief." The complaint must give the defendant "fair notice of what the claim is and the grounds upon which it rests." Bell Atlantic Corp. v. Twombly, 550 U.S. 544, 555 (2007). To meet this requirement, the complaint must be supported by factual allegations. Ashcroft v. Iqbal, 129 S. Ct. 1937, 1950 (2009). "While legal conclusions can provide the framework of a complaint," neither legal conclusions nor conclusory statements are themselves sufficient, and such statements are not entitled to a presumption of truth. Id. at 1949-50.

Under Federal Rule of Civil Procedure 12(b)(6), a complaint may be dismissed for failure to state a claim on which relief can be granted. A complaint may fail to show a right to relief either by lacking a cognizable legal theory or by lacking sufficient facts alleged under a cognizable legal theory. Balistreri v. Pacifica Police Dept., 901 F.2d 696, 699 (9th Cir. 1990).

³The Court applies the same standards it applied on Oracle's previous motion to dismiss and strike, which is repeated here for the convenience of the reader.

In order to survive a motion to dismiss under Rule 12(b)(6), a complaint must "contain either
direct or inferential allegations respecting all the material elements necessary to sustain recovery
under some viable legal theory." Twombly, 550 U.S. at 562 (quoting Car Carriers, Inc. v. Ford
Motor Co., 745 F.2d 1101, 1106 (7th Cir. 1984)) (internal quotations omitted; emphasis in original).
Together, <i>Iqbal</i> and <i>Twombly</i> represent "a two-step process for evaluation of motions to dismiss.
The court first identifies the non-conclusory factual allegations, and the court then determines
whether these allegations, taken as true and construed in the light most favorable to the plaintiff,
'plausibly give rise to an entitlement to relief.'" Fallcochia v. Saxon Mortg., Inc., 709 F. Supp. 2d
860, 865 (E.D. Cal. 2010) (citing <i>Iqbal</i> , 129 S. Ct. at 1950; <i>Erickson v. Pardus</i> , 551 U.S. 89 (2007))
Plausibility, as used in Twombly and Iqbal, refers to whether the non-conclusory factual allegations,
when assumed to be true, "allow the court to draw the reasonable inference that the defendant is
liable for the misconduct alleged." Iqbal, 129 S. Ct. At 1949. "The plausibility standard is not akin
to a 'probability requirement,' but it asks for more than a sheer possibility that a defendant has acted
unlawfully." Id. (quoting Twombly, 550 U.S. at 557).

Where a court dismisses for failure to state a claim pursuant to Rule 12(b)(6), it "should grant leave to amend . . . unless it determines that the pleading could not possibly be cured by the allegation of other facts." *Cook, Perkiss & Liehe v. N. Cal. Collection Serv.*, 911 F.2d 242, 247 (9th Cir. 1990).

2. Motion to Strike

Under Rule 12(f) of the Federal Rules of Civil Procedure, the court may strike from any pleading "any insufficient defense or any redundant, immaterial, impertinent or scandalous matter." "The function of a 12(f) motion to strike is to avoid the expenditure of time and money that must arise from litigating spurious issues by dispensing with those issues prior to trial . . ." Whittlestone, Inc. v. Handi-Craft Co., 618 F.3d 970, 973 (9th Cir. 2010) (internal quotations omitted; citations omitted). However, motions to strike are generally disfavored. 5C Charles Alan Wright & Arthur R. Miller, Federal Practice and Procedure § 1380 (3d ed. 2004).

For the Northern District of California

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В. DrugLogic's Inequitable Conduct Affirmative Defense and Counterclaim

1. Inequitable Conduct Under Exergen⁴

To state a claim for inequitable conduct, a party must allege that "(1) an individual associated with the filing and prosecution of a patent application made an affirmative misrepresentation of a material fact, failed to disclose material information, or submitted false material information; and (2) the individual did so with a specific intent to deceive the PTO." Exergen, 575 F.3d at 1327 fn. 3. See also Star Scientific, Inc. v. R.J. Reynolds Tobacco Co., 537 F.3d 1357, 1365 (Fed. Cir. 2008).

Under Exergen, a party must plead inequitable conduct with particularity under Federal Rule of Civil Procedure 9(b). A pleading that merely recites the substantive elements of inequitable conduct, without providing the specific factual bases for the allegations, is inadequate under Rule 9(b). Exergen, 575 F.3d at 1326-27. Thus, to plead inequitable conduct, a party must identify the specific who, what, when, where, and how of the material misrepresentation or omission committed before the patent examiner. Id. at 1327. Although knowledge and intent may be alleged more generally, a party must still allege sufficient facts to justify an inference that a specific individual had knowledge of the material information withheld or the falsity of the material misrepresentation and withheld or misrepresented that information with the intent to deceive. Fed. R. Civ. P. 9(b); Exergen, 575 F.3d at 1328-29.

In Exergen, the court applied this standard and held that the defendant failed to plead inequitable conduct with sufficient particularity. *Id.* at 1329. The court noted three factual deficiencies in the defendant's proposed pleading. *Id.* First, the pleading failed to name the specific individual who knew of the material information withheld from the PTO and withheld it deliberately, instead referring generally to "Exergen, its agents and/or attorneys." *Id.* Second, the pleading did not identify the specific claims, and which limitations in those claims, to which the withheld references were relevant and where in those references material information could be found. Id. The defendant therefore failed to plead the "what" and "where" of material information

⁴For the convenience of the reader, the Court repeats here the pleading requirements for inequitable conduct under *Exergen*, which are also set forth in the Court's August 8 Order.

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allegedly withheld from the PTO. *Id.* Finally, the pleading failed to identify the particular claim limitations that were allegedly missing from the information of record. *Id.* The defendant's allegations that the withheld references were "material" and "not cumulative to the information already of record" were insufficient to explain "why" the information withheld from the PTO was material and not cumulative and "how" an examiner would have used the information withheld. Id. at 1329-30.

The court in Exergen also held that the defendant failed to adequately plead underlying facts to support an inference of knowledge of the material information withheld and the specific intent to deceive the PTO. Id. at 1330. Although the defendant alleged that Exergen was aware of the withheld patent references, it failed to allege a factual basis to infer that a specific individual associated with the prosecution of the patent knew of the specific information in those references allegedly material to the claims of the patent at issue. *Id.* The court explained that because references can be quite long, "one cannot assume that an individual, who generally knew that a reference existed, also knew of the specific material information contained in that reference." Id. In addition, the court held that the defendant's allegation of deceptive intent "on information and belief" was insufficient. Id. Although a party may plead on "information and belief" under Rule 9(b), the party must still put forth sufficient facts upon which the belief is reasonably based. *Id.* The defendant did not provide any information or plausible reason for its belief, and the circumstances alleged did not suggest a "deliberate decision to withhold a known material reference," a "necessary predicate for inferring deceptive intent." *Id.* at 1330-31 (internal quotations omitted; citation omitted).

2. Whether DrugLogic States a Claim for Inequitable Conduct Under the Exergen Standard in the Second Amended Answer

Oracle argues that the Court should strike DrugLogic's Third Affirmative Defense and Dismiss its Fifth Counterclaim because DrugLogic has failed to plead inequitable conduct with sufficient particularity. Based on the standard set forth in *Exergen*, the Court agrees.

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In its August 8 Order, the Court held that: 1) DrugLogic had not adequately alleged the "what" and "where" of Oracle's alleged material omission; and 2) if DrugLogic, in its amended answer and counterclaims adequately alleged that the patent applicants knew about and withheld certain specific information about the allegedly withheld prior art, then they would also have adequately pled deceptive intent. DrugLogic's amended allegations of inequitable conduct are inadequate because DrugLogic has not identified any specific information that was not disclosed in the specification. Rather, the applicants disclosed all four of the claim elements to which the prior art is relevant in the two paragraphs of the specification that address this prior art, as well as the accompanying figures. See '221 Patent, co. 4, lines 2-24 & Figs. 1, 2. Under Exergen, the court may infer deceptive intent when the facts alleged suggest a "deliberate decision" to withhold known material information. Exergen, 575 F.3d at 1331. Here, the allegations do not support an inference that the applicants deliberately decided to withhold material because Oracle specifically disclosed in the specification not only the existence of the prior art references at issue but also that these references were hierarchical relational medical thesauruses; nor has DrugLogic identified any specific information in these references that was not disclosed. See Fiskars, Inc. v. Hunt MFG. Co., 221 F. 3d 1318, 1327 (Fed. Cir. 2000) (affirming district court's dismissal of inequitable conduct counterclaim following bench trial on basis that allegedly withheld information had been disclosed to examiner and stating that "[a]n applicant can not be guilty of inequitable conduct if the reference was cited to the examiner, whether or not it was a ground of rejection by the examiner"); Chip-Mender, Inc. v. Sherwin-Williams Co., 2006 WL 13058, at *7 (N.D. Cal. Jan. 3, 2006) (holding that defendant's counterclaim for Walker Process fraud based on allegation that patent applicant disclosed prior art reference to patent examiner but did not disclose certain details about that reference failed to state a claim because allegations did not support inference of intent to deceive); Transclean Corp. v. Bridgewood Servs., Inc., 2000 WL 33175724, at *8 (D. Minn. Nov. 1, 2008) (holding after a bench trial that inequitable conduct counterclaim failed because allegedly withheld information had been disclosed in patent application and rejecting as "implausible, if not perverse" the defendant's contention that the applicants had only referenced the allegedly withheld prior art in

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their application to "shield the substantive pertinence of that [prior art] from the Examiner's attention").

Accordingly, DrugLogic's inequitable conduct counterclaim is dismissed with prejudice under Rule 12(b)(6). The Court strikes under Rule 12(f) DrugLogic's inequitable conduct affirmative defense.

C. DrugLogic's Unfair Competition Counterclaim and Availability of **Compensatory Damages on that Claim**

Oracle seeks dismissal of DrugLogic's Unfair Competition counterclaim to the extent it is based on common law, citing the California Supreme Court's Bank of the West decision for the proposition that under common law, "unfair competition" is limited to passing off. The Court agrees.

The Court notes at the outset that Bank of the West addresses the meaning of common law unfair competition under California law in the context of an insurance coverage dispute, as do many of the cases that address the nature of common law unfair competition. In particular, in Bank of the West, the court addressed whether a general liability insurance policy providing coverage for damages arising out of unfair competition covered damages that were paid by Bank of the West in an action that was based on alleged unfair business practices in connection with a loan program. 2 Cal. 4th at 1258-1259. The court reasoned that because damages are not available for statutory unfair competition under Cal. Bus. & Prof. Code § 17203, the provision provided coverage only for common law unfair competition, on which damages are available. Id. at 1265-1266. In that context, the court stated as follows:

The common law tort of unfair competition is generally thought to be synonymous with the act of "passing off" one's goods as those of another. The tort developed as an equitable remedy against the wrongful exploitation of trade names and common law trademarks that were not otherwise entitled to legal protection. (See generally 1 Callmann, Unfair Competition Trademarks & Monopolies (4th ed. 1981) §§ 2.01–2.03.) According to some authorities, the tort also includes acts analogous to "passing off," such as the sale of confusingly similar products, by which a person exploits a competitor's reputation in the market. (See Rest., Torts, §§ 711–743; see also 1 Callmann, supra, § 2.04.)

Id. at 1263.

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While arguably dicta, the California Supreme Court's characterization of common law unfair competition in Bank of the West has been followed by the Ninth Circuit in the context of a motion dismiss. See, e.g., Sybersound Records, Inc. v. UAV Corp., 517 F.3d 1137, 1153 (9th Cir. 2008) (affirming dismissal of common law unfair competition claim under Bank of the West on the basis that no passing off was alleged); Southland Sod Farms v. Stover Seed Co. 108 F.3d 1134, 1147 (9th Cir. 1997) (same). These decisions undermine DrugLogic's assertion that the Court should follow Hewlett-Packard Co. v. Cigna Prop. & Cas. Ins., 1999 U.S. Dist. LEXIS 20655 (N.D. Cal. Aug. 24, 1999), in which the court concluded, in an insurance coverage context, that common law unfair competition is not limited to passing off. In Hewlett Packard, the court addressed whether an insurer had a duty to defend and indemnify under a policy that covered "unfair competition" where the claims for unfair competition in the underlying litigation were not based on allegations of passing off. 1999 U.S. Dist. LEXIS 20655, at * 13 (N.D. Cal. Aug. 24, 1999). Following *Bank of* the West, the court reasoned that coverage extended only to common law unfair competition, which can support a claim for damages. *Id.* However, it rejected as dicta the California Supreme Court's characterization of common law unfair competition as limited to passing off. *Id.* Instead, it addressed how a layperson would understand the term "unfair competition," looking beyond California law to general treatises and the law of other jurisdictions. *Id.* For example, the court quoted the statement in Prosser and Keaton on Torts that "[u]nfair competition . . . can be found when the defendant engages in any conduct that amounts to a recognized tort and when that tort deprives the plaintiff of customers or other prospects." *Id.* (quoting 1013 (5th Ed. 1984)). The Ninth Circuit has not adopted a similar approach in cases decided after the *Hewlett-Packard* case, including in Sybersound Records, cited above. Therefore, the Court concludes that DrugLogic fails to state a claim for common law unfair competition.

The Ninth Circuit's decision in *Duncan v. Stuetzle* does not stand for a contrary result. In that case, the complaint was filed in California state court and asserted claims for "1) misappropriation of proprietary information, 2) unfair competition, and 3) civil conspiracy." 76 F.3d at 1483-1484. The complaint did not specify whether the claims were being asserted under state or

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federal law. Id. The defendant removed to federal district court and the court had to determine whether any of the claims were federal claims for the purposes of determining whether removal was proper. Id. at 1484. The district court found that the complaint stated a claim under the Lanham Act and concluded on that basis that federal jurisdiction existed. *Id.* On appeal, the Ninth Circuit held that the district court had erred, concluding that each of the claims asserted – including the claim for unfair competition – existed under California state law. *Id.* at 1486. In that context, the court held that the plaintiff's claim for unfair competition was not a federal claim because "California provides both statutory and common law causes of action for unfair competition." Id. at 1489. The court further held that to the extent the plaintiff requested monetary damages, which are not available under the UCL, this did not mean that the claim was a federal Lanham Act claim (which allows for damages) because damages are also available on a California common law claim for unfair competition. *Id.* at 1489.

The holding of *Duncan v. Stuezle* does not support DrugLogic's position for two reasons. First, the court did not address whether common law unfair competition, under Bank of the West, is limited to passing off; nor did it address whether the plaintiff's allegations were consistent with the characterization of common law unfair competition claims under California law in Bank of the West. See 76 F.3d at 1489-1490. Rather, the court merely stated that "[e]ach of these causes of actions [statutory and common law unfair competition] provides a theoretical state law basis for Duncan's requested relief." Id. at 1489 (emphasis added). Second, the allegations in Duncan v. Stuezle suggest that the unfair competition claim in that case, though not expressly referred to as "passing off," did in fact fall within the scope of common law unfair competition as set forth in Bank of the West. In particular, the plaintiff in *Duncan v. Stuezle* included the following allegations in support of her unfair competition claim:

20. On or about the month of January, 1990, in the County of San Luis Obispo, the Defendants obtained proprietary information regarding the "Footsie Wootsie" foot massage chair, regarding, without limitation, the product's exterior and mechanical design, the Plaintiffs' company's marketing strategy, the product's recent income, and the methods by which the machine was produced. The Defendants afterward began to produce their own foot massage chairs which closely resemble the chairs produced by Plaintiffs, and do not carry any identifying labels or machine numbers to distinguish them from Plaintiffs' product.

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Subsequently, the Defendants began manufacturing and distributing these duplica	tions or
reproductions of the "Footsie Wootsie" foot massage chair throughout, as far as is	s known a
present, the Southern California area.	

21. The foot massage chairs manufactured, distributed, and sold by the Defendants are designed and calculated to deceive and mislead purchasers and consumers of Plaintiffs' foot massage chair. Further, the Defendants' produce has actually deceived, and continues to deceive consumers, and caused them to use the chairs sold by the Defendants, believing that the chairs were manufactured, sold and distributed by the Plaintiffs.

Id. at 1484. Bank of the West recognized that a common law unfair competition claim may include claims such as the one asserted in *Duncan v. Stuezle*, namely, claims that are "analogous to 'passing off,' such as the sale of confusingly similar products, by which a person exploits a competitor's reputation in the market." 2 Cal. 4th at 1263. In contrast, no such analogous claim is alleged here.

Accordingly, the Court concludes that DrugLogic's claim for unfair competition fails to state a claim because no passing off, or any analogous claim, is alleged. Further, because it is undisputed that a statutory unfair competition claim under the UCL cannot give rise to compensatory damages, the Court strikes DrugLogic's request for disgorgement of profits on Counterclaim Six.

Availability of Injunctive Relief on Breach of Contract Claim D.

Oracle asserts that DrugLogic's request for injunctive relief on its breach of contract counterclaim should be stricken, citing to the elements required under California law for specific performance that are set forth in *Tamarind Lithography Workshop*, Inc. v. Sanders, 143 Cal. App. 3d 571, 575 (1983). In *Tamarind Lithography*, the court of appeals held that the trial court had erred in denying a request for specific performance in addition to an award of damages by a jury following a jury trial. 143 Cal. App. 3d at 575. The court stated as follows:

(1) The availability of the remedy of specific performance is premised upon well established requisites. These requisites include: A showing by plaintiff of (1) the inadequacy of his legal remedy; (2) an underlying contract that is both reasonable and supported by adequate consideration; (3) the existence of a mutuality of remedies; (4) contractual terms which are sufficiently definite to enable the court to know what it is to enforce; and (5) a substantial similarity of the requested performance to that promised in the contract.

Id. Oracle contends that DrugLogic has not adequately pled each of these specific elements. It has not, however, cited case law establishing that it is appropriate to strike a request for specific performance at the pleading stage of the case, especially where, as here, a party has included general

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